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⑤4 Process for the production of alkenyl-succinic anhydrides.

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Description

The present invention relates to the preparation of alkenyl-succinic anhydride compounds by the reaction of α -monoolefins with maleic anhydride (MA).

5 Alkenyl-succinic anhydrides are useful industrial compounds. Quoted fields of use are curing agents for synthetic resins, elastomer components, rust and corrosion inhibitors, additives for increasing viscosity in paints and greases, foam suppression, pesticides and fungicides, surfactant additives, plasticizers, intermediates in synthetic chemistry and drug manufacture, etc...

The chemistry of the reaction between MA and α -olefins resulting from the extraction or cracking of 10 natural oils has been well investigated and described in literature and patents.

For instance US 3,855,251 alkenyl succinic anhydrides derived from the non-catalytic addition of maleic anhydride and a butene polymer hydrocarbon.

US-Patents Nos 2,411,215 (KISE et al) and 3,412,111 (IRWIN et al) provide a detailed list of useful operating parameters and conditions pertaining to this reaction. The molar ratio of olefin (C_3 to C_{60} -olefins) 15 to MA can be comprised between about 0.5 to 20 and the reaction can be carried out at temperatures of about 160 °C to about 300 °C for 0.1 to 48 hrs under pressures of about 1 to 1000 lb/square inch (0.069 bar to 69 bar). US 2,411,215 specifies that yields in this reaction are improved if the reaction medium is well homogeneous, i.e. if the MA is completely dissolved in the α -olefins prior to reaction; this can be effected by stirring the ingredients together beforehand at temperatures below 200 °C, the reaction proper being 20 carried out afterwards at temperatures above 200 °C.

Another technique to increase the yields and to reduce the proportion of undesirable polymeric side-products (gums) which sometimes contaminate the desired alkylene-succinic anhydrides is to add polymerization inhibitors to the reaction mixture. This technique is particularly emphasized in US 3,412,111. In 25 this case the preferred inhibitors are aromatic sulfur or hydroxy compounds, e.g. 2,2-bis (p-hydroxyphenyl)-propane, hydroquinone and phenothiazine. US 4,414,397 discloses the preparation of alkenyl substituted aliphatic decarboxylic anhydrides from olefin oligomer and unsaturated dicarboxylic acid in the presence of a catalyst where the by-product production is minimised by the controlled addition of the unsaturated aliphatic dicarboxylic acid anhydride.

At the end of the reaction period, the reaction mixture can be treated in any suitable manner to recover 30 the individual components therein, e.g. by distillation. US-2,411,215 teaches to first drain the gums which accumulate to the bottom of the reaction vessel, then transfer the liquid phase in a flash evaporator in which unreacted MA and α -olefins will evaporate and be recovered by condensation. The residue from the flash evaporator is then distilled in a still for purifying the desired alkenyl-succinic anhydride compounds from remaining residual by-products.

35 In US-3,412,111, there is recommended to effect distillation of the crude reaction product at temperatures of about 50 °C to about 250 °C under about 10^{-4} to about 15 lb/square inch (5×10^{-3} - 760 Torr) to recover unreacted olefin and maleic anhydride (if any present) and purify the desired alkenyl succinic anhydrides.

Although the processes reported in the literature have much merit, it was desirable to still bring 40 improvements thereto, e.g. render the reaction operation simpler, more efficient and more economical, and providing products ready to be used in many instances without further purification by distillation. It was inter alia desirable to decrease as much as possible the production of polymeric side products of the kind disclosed in the prior art. In this connection, it is noted that, even under the optimal disclosed conditions (see US-A-3,412,111) the amount of polymers which had to be separated from the desired product was still 45 12 g in the case of using a mixture of 2 moles of dodecene (337 g) and one mole of MA (98 g) and operating in the presence of phenothiazine. Furthermore, using phenothiazine as an inhibitor was shown to be undesirable by the present inventors as it usually provides a dark colored product.

It was however found quite unexpectedly by the present inventors that using, as the starting α -olefins, a 1:1 to 4:1 by weight mixture, preferably 65:35, of dodecene (D) and tetradecene (T) (C_{12} and C_{14} α -olefins), 50 substantially no polymeric residue was any longer contaminating the desired alkenyl-succinic anhydrides and, as a consequence, the latter can be used in many instances without further purification by distillation. Thus, the method of the present invention which results from this surprising discovery includes the steps of (a) mixing one molar equivalent of maleic anhydride (MA) with α -monoolefins in a quantity corresponding to at least one olefinic bond molar equivalent and heating to a temperature below 200 °C but sufficient to 55 ensure complete dissolution of the MA in the monoolefins and forming a homogeneous solution; (b) heating this homogeneous solution in a closed vessel to a temperature between 200 and 260 °C where the reaction between the MA and the olefin proceeds smoothly to substantially near or full completion relative to the amount of MA used;

(c) distilling off unreacted α -olefins and MA, if any, and isolating the alkenyl-succinic anhydride products for further end-uses, which comprises using as the starting α -monoolefins a 1:1 to 4:1 by weight mixture of dodecene and tetradecene, whereby no significant formation of polymeric by-products is observed, and isolating the desired alkenyl-succinic anhydride products by simply collecting the residue of distillation of step (c), no further purification being needed.

The reasons why this progress arose has not been elucidated. It is however supposed that a factor is the efficient solubility of maleic anhydride in the aforementioned DT mixture, particularly 65:35 by weight, at temperatures below 200 °C, this effect leading to the formation of substantially well homogeneous solutions of the reactants before reaction.

Another factor of improvement is the efficient control of the reaction time in view of the fact that undesirable side products probably form at the end of the reaction between small quantities of unreacted MA and the desired alkenyl-succinic anhydrides. It may therefore be advantageous to stop the reaction at a time before it goes to full completion relative to the MA used, e.g. when only 80 to 95% of the MA has been consumed, and then proceed with the recovery of unreacted materials by reduced pressure distillation. The somewhat reduced yield of alkenyl-succinic anhydride experienced then is entirely compensated by the clean recovery of the unused MA and α -olefins which can be, of course, recycled. It is noted in this connection, that in the case of operating with batches of about 1 to 5 kg of starting materials reacted at 220-240 °C, the preferred reaction times are in the order of 1-10 hrs, more preferably 2-3 hrs.

The polymerization inhibitors to be used in this reaction can be those reported in US-3,412,111 incorporated herein by reference. Phenothiazine for instance is effective but, as said before, provides colored products; consequently non-sulfur inhibitors like hydroquinone are preferred. Excellent results are recorded with mono-etherified hydroquinone, e.g. hydroquinone-lower alkoxy-monoethers such as the monomethyl- or ethyl hydroquinones.

In general the following batch procedure can be carried out to embody the method of the invention.

The 1:1 to 4:1, by weight mixture of D and T, preferably 65:35 by weight (1-5 molar equivalents, preferably 1-2 equiv.) is placed in an autoclave with 0.1 to 2% by weight of inhibitor (calculated on the MA) and heated progressively under a blanket of inert gas (e.g. N₂, CO₂ or argon) to a temperature of about 160 - 200 °C, preferably 170 - 180 °C, and the MA (1 molar equivalent) is added in liquid (molten) form under stirring. At these temperatures, as said before, the solubility of the MA in the α -olefins mixture is satisfactory; separate experiments in the present work have shown that it is about 20-25% by weight at 160 - 180 °C. When the dissolution is complete and the solution is homogeneous, the vessel is sealingly closed and the temperature is raised to a value above 200 °C, preferably 220 - 250 °C, for a period of time of 1 - 10 hrs, preferably 2 - 3 hrs, depending on the desired degree of reaction completion. During the reaction a pressure rise is experienced of about 2 - 2.5 bar (not exceeding 4 - 5 bar). With the longer reaction times, e.g. 5 - 10 hrs, the MA is substantially fully consumed and the amount of it which can be recovered after the end of the reaction is small or negligible. With the shorter reaction times, the amount of unreacted MA is greater but it can be recovered cleanly for recycling at the end of the reaction and the reaction products have a lighter color.

Then, when the reaction is over, heating is discontinued and the reaction mixture is subjected to distillation to recover the unreacted starting materials. During this distillation, it is important that the condenser be kept at the right temperature to avoid plugging by solidified maleic anhydride which solidifies at about 60 °C. An appropriate condenser temperature is about 61 - 70 °C and the pressure during distillation is preferably 12 - 20 Torr, at least at start.

After distillation, the residue is collected and consists of a mixture of dodecenyl- and tetradecenyl-succinic anhydride in proportions substantially equivalent to that in the starting mixture of α -olefins. This reaction product is pure enough for most applications and can be used as such without further purification. The collected recovered α -olefins (the relative proportion of which remains substantially constant) and the MA can then be recycled in a next run without problems since their purity is about equivalent to that of the starting materials.

Although the foregoing general conditions have been described with reference to a batch process, it is obvious that they also apply to a continuous process. In this case, the key components comprise a preliminary mixing chamber into which the α -olefin mixture containing the inhibitor and the molten maleic anhydride are fed simultaneously in a correct mole ratio and of sufficient volume retention time to ensure complete dissolution of the MA in the α -olefins; this preliminary chamber is followed by a line-reactor (e.g. a chamber or a coil) of sufficient length to ensure proper heating of the reactants for the desired time. After reaction, the mixture can be subjected to continuous distillation by usual procedures and using conventional equipment to assure continuous separation of the products and recycling of unused reactants.

The following Examples illustrate the invention in more detail.

Examples 1 to 9

A 65:35 (by weight) mixture of C₁₂ - C₁₄ α -olefins containing 1708 g (10.15 moles) of dodecene (D) and 919 g (4.68 moles) of tetradecene (T) and 7.3 g of hydroquinone was heated to 170°C under nitrogen in a 5 liter autoclave. 726 g (7.40 moles) of molten (100°C) maleic anhydride (MA) were added under stirring and the vessel was tightly closed. After the solution had become homogeneous (192°C), the temperature was raised to 220°C and maintained there for 3 hrs.

The reaction mixture was distilled in a "Rotavapor" (BUCHI, type EL-131) under reduced pressure starting under 14 Torr and ending under 1.6 Torr (bath temperature 150°C). Under these conditions, the approximate boiling temperatures (14 Torr) of the components are: MA 80°C; D 100°C; T 125°C. The condenser was kept at 63°C (thermostated water circulation) to prevent plugging of the apparatus by distilled MA. After distillation, the light colored residue 1940 g (yield 95%) was shown by analysis to be a nearly pure mixture of C₁₂-and C₁₄-alkenyl-succinic anhydrides (DTSA) in a molar ratio very close to that (10.2/4.7) of the initial olefins. The distillate (1389 g) separated into a liquid and a solid phase (17.6 g) which was shown later to be maleic anhydride. The liquid phase was the recovered α -olefins mixture, the components being in a ratio similar to that of the starting material. Taking into account that the solubility of MA in the hydrocarbon mixture at room temperature is about 0.3%, the yield of recovered MA is approximately 3%. The yield of recovered olefins which can be directly recycled in a subsequent run approximates the theoretical value.

Other Examples in which the various parameters involved were varied like temperature, reaction time, mole ratio of D+T to MA, nature of inhibitor (always 1% by weight relative to MA) were run similarly and the data are reported in the Table 1 below. In the Table the % recovered D+T mixture is calculated on the theoretically unused portion (excess of D+T) with the exception of Example 6 (stoichiometric amount of reactants) where the quoted figure is the actual value.

TABLE 1

Example No.	Starting materials		Inhibitor	Temp. (°C)	Time (hrs)	Products		
	D + T mixture (moles)	D + T/MA mole ratio				DTSA (%)	D + T (%)	MA (%)
1	14.8	2	none	220	8	96.4	101.5	-
2	14.8	2	PT	220	8	97.7	102.2	-
3	14.8	2	HQ	220	8	98.2	100.8	-
4	14.8		HQME	220	8	96.2	103.0	-
5	13.8	1.5	HQ	220	8	94.6	112.6	-
6	12.2	1	HQ	220	8	88.2	13.5	0.2
7	14.8	2	HQ	220	3	95.1	103.9	3.0
8	13.8	1.5	HQ	240	8	94.8	111.8	-
9	13.8	1.5	HQ	220	2	89.5	119.5	10.1
Codes:								
PT = phenothiazine								
HQ = hydroquinone								
HQME = monomethylether hydroquinone								

Example 10 (1:1 by weight D:T mixture)

A mixture of 1258.2 g (7.48 moles) of dodecene, 1258.2 g (6.41 moles) of tetradecene (total 13.88 moles), 902 g (9.19 moles) of maleic anhydride and 9 g of hydroquinone was stirred and heated to 190°C in a pressure reactor as in the previous examples until homogeneous.

Then the temperature was raised to 220°C under sealed conditions, whereby the reaction started. The temperature was maintained and samples were taken at intervals of time through a bottom outlet valve of the reactor and analyzed by usual means for concentration of constituents (see thereafter). The results are given in Table 2 in % by weight up to a period of 5 hrs.

TABLE 2

Time (hrs)	MA	D	T	DTSA	
				12	14
0	26.3	36.7	36.7	0	0
1/2	11.6	18.8	18.4	14.9	15.7
1	7.14	17.1	17.9	19.2	20.2
2	3.21	14.4	14.1	21.0	21.8
3	1.38	13.5	13.7	22.5	23.1
4	0.60	13.6	13.6	24.2	24.6
5	0.28	14.9	14.0	25.5	25.9

After 5 hrs, the reaction was stopped and the mixture (3409 g) subjected to vacuum distillation (bath 150° C) as indicated in the previous examples; a total of 896 g (4.94 moles) of unreacted D and T olefins of weight ratio substantially near 1:1 was collected, whereby the yield of DTSA based on used up olefins (i.e. 8.94 moles) and the residue weight (2496 g \approx 8.9 moles) was about 99%.

Example 11 (4:1 by weight D/T mixture)

The reaction conditions and subsequent manipulations were repeated exactly as in Example 10 using the following components and conditions:

Dodecene	2013.1 g (11.96 moles)
Tetradecene	503.3 g (2.55 moles)
Maleic anhydride	902.1 g (9.2 moles)
Hydroquinone	9 g
D + T / MA ratio	(1.58 mole/mole)
Reaction time	5 hrs
Reaction temperature	220° C
Distillation of reaction mixture	150° C/15-4 Torr

Analyses performed as in the previous example are reported in Table 3 below.

TABLE 3

Time (hrs)	MA	D	T	DTSA	
				12	14
0	26.3	58.7	14.7	0	0
1/2	12.0	37.2	10.7	25.3	6.6
1	7.51	30.0	8.5	29.2	8.6
2	2.62	21.8	5.6	38.6	9.8
3	1.45	21.9	5.6	41.9	10.5
4	0.53	22.5	6.2	44.3	13.3
5	0.22	21.7	6.1	42.5	11.7

Distillation provided 9.83 g (5.6 moles) of olefins and 2421 g (8.9 moles) of DTSA. Hence the overall yield of DTSA was about 99%.

Analyses

The reaction mixture was analyzed by sampling through a bottom outlet valve of the reactor and subjecting to GLC.

The gas-liquid chromatograph was a "CARLO ERBA Fractovap Series 4160" equipped with a 30 m x 0.32 mm J & W fused silica capillary column (liquid phase: 0.25 μ m DB-5).

5	- carrier gas	H ₂ , 1.5 ml/min, inlet pressure 0.4 atm.
	- detector	FID, range 10 mv, attenuation 8, temperature 250 ° C.
	- injection	1-1.5 μ l, split ~ 1/9, temperature 230 ° C.
	- column temperature	initial: 80 ° C for 2 min, final: 240 ° C for 7 min, heating rate 10 ° C/min.

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Claims

1. A method for manufacturing alkenyl-succinic anhydrides in a form sufficiently pure for using them directly, this method including the steps of:
 - (a) mixing one molar equivalent of maleic anhydride (MA) with α -monoolefins in a quantity corresponding to at least one olefinic bond molar equivalent and heating to a temperature below 200 ° C but sufficient to ensure complete dissolution of the MA in the monoolefins and forming a homogeneous solution;
 - (b) heating this homogeneous solution in a closed vessel to a temperature between 200 and 260 ° C where the reaction between the MA and the olefin proceeds smoothly to substantially near or full completion relative to the amount of MA used;
 - (c) distilling off unreacted α -olefins and MA, if any, and isolating the alkenyl-succinic anhydride products for further end-uses,
- 25 which comprises using as the starting α -monoolefins a 1:1 to 4:1 by weight mixture of dodecene and tetradecene, whereby no significant formation of polymeric by-products is observed, and isolating the desired alkenyl-succinic anhydride products by simply collecting the residue of distillation of step (c), no further purification being needed.
- 30 2. The method of claim 1, wherein dissolution in step (a) is performed at 160 - 180 ° C and step (b) is performed for 0.2 to 10 hrs at 220 - 240 ° C.
3. The method of claim 2, wherein step (b) is carried out for 2 - 3 hrs and the amount of consumed MA is 80 - 95% by weight.
- 35 4. The method of claim 1, which comprises adding 0.1 - 2% of a polymerization inhibitor to the reaction solution.
5. The method of claim 4, wherein the inhibitor is selected from at least one of hydroquinone, monoethers of hydroquinone and phenothiazine.
- 40 6. The method of claim 1, which comprises recycling the unreacted α -olefins and/or MA recovered from step (c).
- 45 7. The method of claim 1, wherein the distillation in step (c) is effected under 12 - 20 Torr, the condenser temperature being controlled to a value above the melting temperature of MA.
8. The method of claim 1, wherein the starting mono- α -olefins mixture is a 65:35 by weight mixture of dodecene and tetradecene.

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Patentansprüche

1. Verfahren zur Herstellung von Alkenylbernsteinsäureanhydriden in einer für deren direkte Verwendung ausreichend reinen Form, welches Verfahren die folgenden Schritte umfaßt:
 - (a) Mischen eines Moläquivalents Maleinsäureanhydrid (MA) mit α -Monoolefinen in einer Menge entsprechend mindestens einem Moläquivalent einer olefinischen Bindung und Erwärmen auf eine Temperatur unterhalb 200 ° C, die jedoch ausreichend ist, um eine vollständige Auflösung des MA in den Monoolefinen sicherzustellen, und Bilden einer homogenen Lösung;

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(b) Erwärmen dieser homogenen Lösung in einem geschlossenen Behälter auf eine Temperatur zwischen 200 und 260 °C, bei welcher die Umsetzung zwischen dem MA und dem Olefin glatt voranschreitet bis im wesentlichen nahe der oder der vollständigen Umsetzung bezüglich der Menge des verwendeten MA;

5 (c) Abdestillieren von nichtumgesetzten α -Olefinen und MA, falls vorhanden, und Isolieren der Alkenylbernsteinsäureanhydridprodukte für weitere Endanwendungen.

umfassend die Verwendung einer Mischung aus Dodecen und Tetradecen im Gewichtsverhältnis von 1:1 bis 4:1 als Ausgangs- α -Monoolefine, wodurch keine signifikante Bildung von polymeren Nebenprodukten erhalten wird, und die Isolierung der erwünschten Alkenylbernsteinsäureanhydridprodukte durch einfaches Sammeln des Rückstandes der Destillation des Schritts (c), wobei keine weitere Reinigung erforderlich ist.

2. Verfahren nach Anspruch 1, wobei die Auflösung in Schritt (a) bei 160-180 °C durchgeführt und Schritt (b) während 0,2 bis 10 Stunden bei 220-240 °C durchgeführt wird.

15 3. Verfahren nach Anspruch 2, wobei Schritt (b) während 2-3 Stunden durchgeführt wird und die Menge an verbrauchtem MA 80-95 Gew.-% beträgt.

4. Verfahren nach Anspruch 1, umfassend die Zugabe von 0,1-2 % eines Polymerisationsinhibitors zu der Reaktionslösung.

5. Verfahren nach Anspruch 4, wobei der Inhibitor aus mindestens einem aus Hydrochinon, Monoethern von Hydrochinon und Phenothiazin gewählt wird.

25 6. Verfahren nach Anspruch 1, umfassend die Wiederverwendung der aus Schritt (c) rückgewonnenen nichtreagierten α -Olefine und/oder MA.

7. Verfahren nach Anspruch 1, wobei die Destillation in Schritt (c) unter 12-20 Torr bewirkt wird, und wobei die Kühlertemperatur auf einen Wert oberhalb der Schmelztemperatur von MA reguliert wird.

30 8. Verfahren nach Anspruch 1, wobei die Ausgangsmischung der α -Monoolefine eine Mischung aus Dodecen und Tetradecen im Gewichtsverhältnis von 65:35 ist.

Revendications

35 1. Procédé pour la fabrication d'anhydrides alcénylsucciniques sous une forme suffisamment pure pour les utiliser directement, ce procédé comprenant les étapes consistant:

(a) à mélanger un équivalent molaire d'anhydride maléique (MA) avec des α -monoooléfines en une quantité correspondant à au moins un équivalent molaire de liaison oléfinique et à chauffer à une température inférieure à 200 °C mais suffisante pour assurer la dissolution complète du MA dans les monooléfines et à former une solution homogène;

(b) à chauffer cette solution homogène dans un récipient fermé à une température comprise entre 200 et 260 °C, où la réaction entre le MA et l'oléfine progresse régulièrement vers l'achèvement total ou sensiblement total par rapport à la quantité de MA utilisée;

45 (c) à retirer par distillation les α -oléfines n'ayant pas réagi et MA, s'il y en a, et à isoler les produits anhydride alcényl-succinique pour d'autres utilisations finales,

qui consiste à utiliser, comme α -monoooléfines de départ, un mélange de 1:1 à 4:1 en poids de dodécène et de tétradécène, ainsi on n'observe aucune formation significative de sous-produits polymères, et à isoler les produits anhydride alcényl-succinique souhaités en recueillant simplement le résidu de distillation de l'étape (c), aucune purification supplémentaire n'étant nécessaire.

2. Procédé selon la revendication 1, dans lequel la dissolution dans l'étape (a) est réalisée à une température de 160 à 180 °C et l'étape (b) est réalisée pendant 0,2 à 10 heures à une température de 220 à 240 °C.

55 3. Procédé selon la revendication 2, dans lequel l'étape (b) est effectuée pendant 2 à 3 heures et la quantité de MA consommée est comprise entre 80 et 95% en poids.

4. Procédé selon la revendication 1, qui consiste à ajouter 0,1 à 2% d'un inhibiteur de polymérisation à la solution réactionnelle.
5. Procédé selon la revendication 4, dans lequel l'inhibiteur est choisi parmi au moins un de l'hydroquinone, des mono-éthers d'hydroquinone et de la phénothiazine.
6. Procédé selon la revendication 1, qui comprend le recyclage des α -oléfines n'ayant pas réagi et/ou du MA récupérés à partir de l'étape (c).
- 10 7. Procédé selon la revendication 1, dans lequel la distillation dans l'étape (c) est effectuée sous une pression de 1,6 à 2,7 kPa (12 à 20 torr), la température du réfrigérant étant réglée à une valeur au-dessus de la température de fusion du MA.
- 15 8. Procédé selon la revendication 1, dans lequel le mélange de mono- α -oléfines de départ est un mélange de 65:35 en poids de dodécène et de tétradécène.

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